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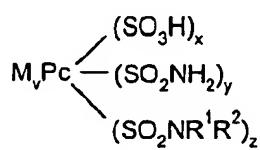
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#### (54) Compounds, compositions and use

(57) Phthalocyanine compounds of Formula (1) have utility as colorants in inks for ink-jet printing, where Formula (1) represents:



### Formula (1)

R<sup>3</sup> is independently a group selected from H, -SR<sup>4</sup>, -CO<sub>2</sub>R<sup>5</sup> and -NR<sup>6</sup>R<sup>7</sup>; where:  
 R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup> and R<sup>7</sup> independently represent H, C<sub>1-30</sub>alkyl optionally substituted by one or more groups selected from hydroxy, mercapto, sulpho, carboxy, cyano and -PO<sub>3</sub>H<sub>2</sub>;  
 v is the valence of P<sub>c</sub> divided by the valence of M;  
 x is from 1.8 to 3.8;  
 y is from 0.1 to 2.7;  
 z is from 0.1 to 2.7; and  
 x, y and z satisfy 2 ≤ x + y + z ≤ 4.

in which:

M represents a metal or H;  
 Pc represents a phthalocyanine nucleus; and  
 R<sup>1</sup> represents H or -(CH<sub>2</sub>)<sub>n</sub>R<sup>3</sup>; R<sup>2</sup> represents  
 -(CH<sub>2</sub>)<sub>n</sub>R<sup>3</sup>; or R<sup>1</sup> and R<sup>2</sup> together with the nitrogen  
 atom to which they are attached represent a 5 or  
 6-membered ring; where: n is independently an in-  
 teger from 1 to 30; and

### Description

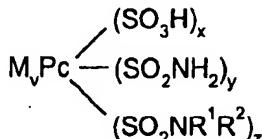
[0001] This invention relates to compounds, to ink compositions comprising them and to the use of such inks in ink jet printing ("IJP"). IJP is a non-impact printing technique in which droplets of ink are ejected through a fine nozzle onto a substrate without bringing the nozzle into contact with the substrate.

[0002] There are many demanding performance requirements for colorants and inks used in IJP. For example they desirably provide sharp, non-feathered images having good water-fastness, light-fastness and optical density. The inks are often required to dry quickly when applied to a substrate to prevent smudging, but they should not form a crust over the tip of an ink jet nozzle because this will stop the printer from working. The inks should also be stable to storage over time without decomposing or forming a precipitate which could block the fine nozzle.

[0003] It is essential that colorants used in IJP have good solubility (preferably in aqueous systems) so they can be dissolved in ink and fired from the Ink-jet head without coming out of solution and blocking the nozzle. Yet colorants that have good operability (i.e. high water solubility) are likely to produce prints of poor water fastness due to the same high water solubility. A significant challenge in designing new colorants for IJP is to provide in the same molecule the apparently mutually exclusive properties of good operability for the ink and yet high water fastness for the print. This often requires the synthesis of a colorant molecule of complex structure comprising many different functional groups and such complex molecules can be expensive and difficult to prepare in high yield.

[0004] It would be advantageous to provide colorants which overcome some or all of the preceding disadvantages.

[0005] Therefore according to the present invention there are provided one or more compounds of Formula (1) and salts thereof:



### Formula (1)

in which;

M represents a metal or H;

Pc represents a phthalocyanine nucleus; and

R<sup>1</sup> represents H or -(CH<sub>2</sub>)<sub>n</sub>R<sup>3</sup>; R<sup>2</sup> represents (CH<sub>2</sub>)<sub>n</sub>R<sup>3</sup>; or R<sup>1</sup> and R<sup>2</sup> together with the nitrogen atom to which they are attached represent a 5 or 6-membered ring; where:

**n** is independently an integer from 1 to 30; and

$R^3$  is independently a group selected from H, hydroxy, sulpho, cyano,  $-SR^4$ ,  $-CO_2R^5$ ,  $-PO_3H_2$  and  $-NR^6R^7$ ; where:  $R^4$ ,  $R^5$ ,  $R^6$  and  $R^7$  independently represent H,  $C_{1-6}$ alkyl optionally substituted by one or more groups selected

from hydroxy, mercapto, sulpho, carboxy, cyano and  $\text{-PO}_3\text{H}_2$ .

v is the valence of  $P_c$  divided by the valence of  $M$ :

$x$  is from 1.2 to 3.8;

v is from 0.1 to 2.7;

$z$  is from 0.1 to 2.7; and

x, y and z satisfy  $2 \leq x + y + z \leq 4$ :

with the provisos that:

50 R<sup>4</sup> and R<sup>5</sup> are other than H;  
 when R<sup>1</sup> is H or C<sub>1-4</sub>alkyl then R<sup>4</sup> is other than C<sub>1-4</sub>alkyleneSO<sub>3</sub>H;  
 when R<sup>1</sup> is H or C<sub>1-4</sub>alkyl and R<sup>6</sup> is H, C<sub>1-4</sub>alkyl or hydroxyC<sub>1-4</sub>alkyl; then R<sup>7</sup> is other than C<sub>1-4</sub>alkyl, hydroxyC<sub>1-4</sub>alkyl or morpholino; and  
 55 when R<sup>1</sup> is H or C<sub>1-4</sub>alkyl then R<sup>6</sup> and R<sup>7</sup> are other than both H or (together with the N atom to which they are attached) other than a morpholino ring.

[0006] The applicant has discovered that the above phthalocyanine compounds have particular utility as colorants for use in IJP. Compared to analogues with one or no sulphonic acid groups, the applicant has found compounds of

the present invention exhibit improved operational advantages when used in IJP without adversely effecting print water fastness. In addition, compounds of the present invention have other properties which are also useful in IJP. For example they can produce IJ prints which exhibit good optical density, light fastness and/or a particularly attractive cyan shade. Their preparation is straightforward because of their simpler structure.

5 [0007] The compounds of the present invention described herein include all chemical and physical forms thereof (such as those described herein) and incorporate all the above provisos. Preferred compounds are those which are IJP-effective.

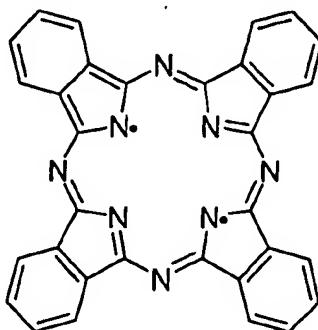
[0008] When M is a metal it is preferably selected from Li, Na, K, Mg, Ca, Ba, Al, Si, Sn, Pb, Rh, Sc, Ti, V, Cr, Mn, Fe, Co, Ni and Cu, more preferably from Sc, Ti, Va, Cr, Mn, Fe, Co, Zn, Ni and Cu; especially from Ni and Cu, particularly 10 Cu.

[0009] It will be appreciated that v (which denotes the stoichiometric ratio of M to Pc) is the molar ratio which produces a stable complex. It can readily be determined from the relative valences of Pc and M (=  $Pc/M$ ). Thus preferably if  $Pc$  is a bivalent radical of formula (2):

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Formula (2)

30 (i.e. the valence of  $Pc$  is 2) then  $v$  is inverse half the valence of M.

[0010] It will also be appreciated that because of the nature of the preparation of compounds of the present invention, in general they may exist as a mixture of different compounds. Thus in Formula (1) the values for  $v$ ,  $x$ ,  $y$  and  $z$  may represent averages for the mixture. Insofar as purification techniques will allow, it is preferred that the mixture comprises one compound as the major component, more preferably the mixture is substantially free of other compounds of Formula (1).

35 [0011] Preferably  $x$  is from 1.2 to 2.8, more preferably 1.5 to 2.5, most preferably 1.8 to 2.2, especially 2.0.

[0012] Preferably  $y$  and  $z$  are independently from 0.3 to 2.0, more preferably from 0.5 to 1.5, most preferably 1.0.

[0013] When  $z$  is  $> 1$  (i.e. where there is a plurality of  $-SO_2NR^1R^2$  groups attached to the  $Pc$  nucleus) each  $R^1$  and  $R^2$  may independently represent a different substituent in each  $-SO_2NR^1R^2$  group.

40 [0014] Most preferred compounds are those in which  $x$  is 2,  $y$  is 1 and  $z$  is 1.

[0015] When  $R^1$  and/or  $R^2$  are  $-(CH_2)_nR^3$  then  $n$  is independently: preferably from 1 to 15, more preferably from 1 to 8, most preferably from 1 to 4.

[0016] When  $R^1$ ,  $R^2$  and the N-atom to which they are attached form a cyclic group, it is preferably selected from morpholino, pyridyl or piperidino; more preferably morpholino.

45 [0017] Preferred compounds of Formula (1) are those in which:

M is Cu or Ni,

50  $R^1$  is selected from H and optionally substituted  $C_{1-15}$ alkyl; and  
 $R^2$  is a substituted  $C_{1-15}$ alkyl.

50

[0018] More preferred compounds of Formula (1) are those in which:

M is Cu,  
 $R^1$  is selected from H,  $C_{1-4}$ alkyl and hydroxy $C_{1-4}$ alkyl; and  
 $R^2$  is a hydroxy $C_{1-4}$ alkyl.

55 [0019] Specific compounds of Formula (1) are selected from those exemplified herein and any IJP-effective salts thereof, more preferably the alkali metal or optionally substituted ammonium salts thereof.

[0020] Any radical group mentioned herein as a substituent refers to a monovalent radical unless otherwise stated. A group which comprises a chain of three or more atoms signifies a group in which the chain may be straight or branched or the chain or any part of the chain may form a ring. Substituents may replace any H attached to an atom in the ring (e.g.  $\text{Pc}$  nucleus) or chain which is chemically suitable and may be located at any available position on the ring or chain (e.g.  $\text{R}^3$  can be on any position on the alkyl chain and is not restricted to terminal position). Preferably the substituents on the  $\text{Pc}$  nucleus are positioned so that none of its phenyl moieties have more than one substituent. The total number of certain atoms is specified herein for certain substituents, for example  $\text{C}_{1-\text{m}}$  alkyl, signifies an alkyl group having from 1 to  $\text{m}$  carbon atoms.

[0021] The terms 'optionally substituted' and 'substituted' as used herein, unless immediately followed by a list of one or more substituent groups, means (optionally) substituted with one or more groups selected from: hydroxy, mercapto, carboxy, sulpho and cyano.

[0022] Unless the context clearly indicates otherwise, as used herein plural forms of the terms herein are to be construed as including the singular form and vice versa.

The term 'IJP effective' (for example with reference to the inks, compositions, ingredients, substituents and/or compounds described herein) will be understood to mean effective for use in ink-jet printing by for example: providing desirable properties to the ink, being compatible with any inert carriers and/or diluents suitable for formulating such inks, being compatible with ink jet printers and/or capable of being ink-jet printed. In relation to the processes described herein effective compounds are those which will undergo the specified reactions to form the compounds of the present invention. Preferably compounds acceptable for use in IJP are Ames negative.

[0023] It will be appreciated compounds of Formula (1) may exist in many different physical and chemical forms which also form part of the present invention. These forms may comprise any of the following (including mixtures thereof and combinations thereof in the same molecular moiety): salts, stereoisomers (e.g. enantiomers, diastereoisomers, geometric isomers, tautomers and/or conformers), zwitterions, polymorphic forms (e.g. phases, crystalline forms, amorphous forms, solid solutions and/or interstitial compounds); complexes (e.g. in addition to the metal  $\text{Pc}$  complexes described herein, chelates, solvates, hydrates and/or complexes with any other suitable ligand) and/or isotopically substituted forms (optionally radio-active, e.g. used as means for selective imaging of the compounds and/or inks containing them and/or as tools to investigate their mode of action in IJP).

[0024] Compounds of Formula (1) may be in the form as shown in the structures herein (i.e. with free sulphonic acid groups) but are preferably in the form of salts. Salts of Formula (1) may be formed from one or more organic and/or inorganic bases and/or acids and compounds of Formula (1) which are acidic and/or basic (for example acid and/or base addition salts). Salts of Formula (1) comprise all IJP-effective salts that may be formed from monovalent and/or multivalent acids and/or bases. Salts of Formula (1) also comprise all enantiomeric salts formed with IJP-effective chiral acids and/or bases and/or any mixtures of enantiomers of such salts (for example racemic mixtures). The dyes may be converted into a salt using known techniques. The present invention comprises all salts of Formula (1) and mixtures thereof, especially those which are IJP-effective.

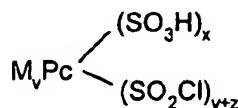
[0025] Preferred salts of Formulae (1) are alkali metal salts (especially lithium, sodium and potassium salts), and optionally substituted ammonium salts (especially salts with ammonia and volatile amines). More preferred salts are those with a cation of formula  $^+\text{NT}_4$  where each T is independently H or optionally substituted alkyl, or two groups represented by T are H or optionally substituted alkyl and the remaining two groups represented by T, together with the N atom to which they are attached, form a 5 or 6 membered ring (preferably a morpholine, pyridine or piperidine ring). Most preferably each T is independently H or  $\text{C}_{1-4}$  alkyl, especially H,  $\text{CH}_3$  or  $\text{CH}_3\text{CH}_2$ , more especially H. Examples of cations which are IJP-effective comprise:  $^+\text{NH}_4$ , morpholinium, piperidinium, pyridinium,  $(\text{CH}_3)_3\text{N}^+\text{H}$ ,  $(\text{CH}_3)_2\text{N}^+\text{H}_2$ ,  $\text{H}_2\text{N}^+(\text{CH}_3)(\text{CH}_2\text{CH}_3)$ ,  $\text{CH}_3\text{N}^+\text{H}_3$ ,  $\text{CH}_3\text{CH}_2\text{N}^+\text{H}_3$ ,  $\text{H}_2\text{N}^+(\text{CH}_2\text{CH}_3)_2$ ,  $\text{CH}_3\text{CH}_2\text{CH}_2\text{N}^+\text{H}_3$ ,  $\text{CH}_3\text{CH}_2\text{CH}_2\text{N}^+\text{H}_3$ ,  $(\text{CH}_3)_2\text{CHN}^+\text{H}_3$ ,  $\text{N}^+(\text{CH}_3)_4$ ,  $\text{N}^+(\text{CH}_2\text{CH}_3)_4$ , N-methyl pyridinium, N,N-dimethyl piperidinium and N,N-dimethyl morpholinium.

[0026] Compounds of Formula (1) may have many uses other than IJP. For example as well as colorants for IJP inks, compounds of the present invention may be used as intermediates in the preparation and/or purification of other compounds of Formula (1) and/or as research tools and/or diagnostic aids in relation to IJP.

[0027] The term 'colorant' as used herein includes both dyes and pigments. Colorants are not limited to materials which solely provide colour in the visible region of the electromagnetic (EM) spectrum but include materials (which may be visibly colourless or weakly coloured) which attenuate radiation in other regions of the EM spectrum invisible to the naked eye [e.g. ultra-violet (UV) and/or infra-red (IR) absorbers]. Compounds of the present invention may exhibit such colorant properties outside the visible region.

[0028] Compounds of Formula (1) may be prepared by the methods described below and by other suitable methods analogous to those described in the art for similar phthalocyanine compounds. A preferred method for preparing one or more compounds of Formula (1) comprises the following steps:

(i) condensing one or more compounds of Formula (3)



Formula(3)

with ammonia and an amine of formula  $\text{NHR}^1\text{R}^2$ ;

10 (ii) treating the adduct from step (i) with a base, to produce one or more compounds of Formula (1);

where  $v$ ,  $x$ ,  $y$ ,  $z$ ,  $M$ ,  $\text{Pc}$ ,  $\text{R}^1$  and  $\text{R}^2$  are as represented herein.

[0029] Preferably in step (i) the ammonia and amine are present respectively in approximately  $y$  and  $z$  molar equivalents to the amount of phthalocyanine.

15 [0030] Preferably in step (ii) the base comprises sodium hydroxide and the pH of the mixture may be adjusted to be from 7.0 to 9.0, preferably 8.0.

20 [0031] Compounds of Formula (3) may be prepared using known methods. A preferred method preparing these compounds comprises heating (preferably for about 1 to about 24 hours), a metal-free or metal containing phthalocyanine (optionally comprising an average of two to four sulpho groups per molecule) with chlorosulphonic acid, preferably at a temperature above 60°C, more preferably above 100°C, most preferably from 120°C to 165°C. Optionally this may be followed cooling, preferably to a temperature from 30°C to 50°C. The phthalocyanine is then heated with  $\text{PCl}_3$  (conveniently for about 4 to about 6 hours), preferably at lower temperature than with the chlorosulphonic acid, more preferably from 80°C to 105°C.

25 [0032] The reactions leading to the formation of the present compounds may be performed under conditions that have been described in the art and compounds of Formula (1) may be isolated by known methods such as spray drying or precipitation followed by filtration.

[0033] In a further aspect of the present Invention there is provided an ink which is effective for use in ink jet printing, the ink comprising a fluid medium, (optionally a liquid) and a colorant (preferably a dye) comprising one or more compounds of the present Invention as defined herein.

30 [0034] Preferably the ink of the present invention comprises:

- (a) from 0.01 to 30 parts of an IJP-effective compound of Formula (1); and
- (b) from 70 to 99.99 parts of a liquid medium or a low melting point solid medium;

35 wherein all parts are by weight and the number of parts of (a)+(b)=100.

[0035] The number of parts of component (a) is preferably from 0.1 to 20, more preferably from 0.5 to 15, and especially from 1 to 5 parts. The number of parts of component (b) is preferably from 99.9 to 80, more preferably from 99.5 to 85, especially from 99 to 95 parts.

40 [0036] When the medium is a liquid, preferably component (a) is completely dissolved in component (b). Preferably component (a) has a solubility in component (b) at 20°C of at least 10%. This allows the preparation of concentrates which may be used to prepare more dilute inks and reduces the chance of the colorant precipitating if evaporation of the liquid medium occurs during storage.

[0037] Preferred liquid media include water, a mixture of water and an organic solvent and an organic solvent free from water.

45 [0038] When the medium comprises a mixture of water and an organic solvent, the weight ratio of water to organic solvent is preferably from 99:1 to 1:99, more preferably from 99:1 to 50:50 and especially from 95:5 to 80:20. Preferably the organic solvent comprising the mixture of water and organic solvent is a water-miscible organic solvent or a mixture of such solvents. The liquid medium may comprise water and preferably two or more, more preferably from 2 to 8, water-soluble organic solvents.

50 [0039] Preferred water-miscible organic solvents comprise:

$\text{C}_{1-6}$ -alkanols, preferably methanol, ethanol, n-propanol, isopropanol, n-butanol, sec-butanol, tert-butanol, n-pentanol, cyclopentanol and/or cyclohexanol;

linear amides, preferably dimethylformamide and/or dimethylacetamide;

55 ketones and/or ketone-alcohols, preferably acetone, methyl ether ketone, cyclohexanone and/or diacetone alcohol; water-miscible ethers, preferably tetrahydrofuran and/or dioxane;

diols, preferably  $\text{C}_{2-12}$ diols (for example pentane-1,5-diol, ethylene glycol, propylene glycol, butylene glycol, pentylene glycol, hexylene glycol and/or thiodiglycol) and/or oligo- and/or poly-alkyleneglycols (for example diethylene

glycol, triethylene glycol, polyethylene glycol and/or polypropylene glycol);  
 5 triols, preferably glycerol and/or 1,2,6-hexanetriol;  
 $C_{1-4}$ alkyl ethers of diols, preferably mono $C_{1-4}$ alkyl ethers of  $C_{2-12}$ diols: {for example 2-methoxyethanol, 2-(2-methoxyethoxy)ethanol, 2-(2-ethoxyethoxy)-ethanol, 2-[2-(2-methoxyethoxy)ethoxy]-ethanol, 2-[2-(2-ethoxyethoxy)ethoxy]-ethanol and/or ethyleneglycol monoallyl ether};  
 10 cyclic amides, preferably 2-pyrrolidone, N-methyl-2-pyrrolidone, N-ethyl-2-pyrrolidone, caprolactam and/or 1,3-dimethylimidazolidone;  
 cyclic esters, preferably caprolactone;  
 sulphoxides, preferably dimethyl sulphoxide and/or sulpholane; and/or  
 10 all IJP-effective mixtures thereof.

**[0040]** More preferred water-soluble organic solvents are selected from:

15 cyclic amides (e.g. 2-pyrrolidone, N-methyl-pyrrolidone and N-ethyl-pyrrolidone);  
 diols, (e.g. 1,5-pentane diol, ethyleneglycol, thiodiglycol, diethyleneglycol and triethyleneglycol);  
 $C_{1-4}$ alkyl ethers of diols (e.g. 2-methoxy-2-ethoxy-2-ethoxyethanol); and all IJP-effective mixtures thereof.

**[0041]** A preferred liquid medium comprises:

20 (a) from 75 to 95 parts water; and  
 (b) from 25 to 5 parts in total of one or more solvents selected from:  
 diethylene glycol, 2-pyrrolidone, thiodiglycol, N-methylpyrrolidone, cyclohexanol, caprolactone, caprolactam  
 25 and pentane-1,5-diol;

where the parts are by weight and the sum of the parts (a) + (b) = 100.

**[0042]** Another preferred liquid medium comprises:

30 (a) from 60 to 80 parts water;  
 (b) from 2 to 20 parts diethylene glycol; and  
 (c) from 0.5 to 20 parts in total of one or more solvents selected from:  
 2-pyrrolidone, N-methylpyrrolidone, cyclohexanol, caprolactone, caprolactam, pentane-1,5-diol and thiodigly-  
 35 col;

where the parts are by weight and the sum of the parts (a) + (b) + (c) = 100.

**[0043]** Examples of further IJP-effective media for inks of the present invention comprise a mixture of water and one or more organic solvents are described in US 4,963,189, US 4,703,113, US 4,626,284 and EP 0425150-A.

**[0044]** When the liquid medium comprises an organic solvent free from water, (i.e. less than 1% water by weight) the solvent preferably has a boiling point of from 30° to 200°C, more preferably of from 40° to 150°C, especially from 50 to 125°C. The organic solvent may be water-immiscible, water-miscible or a mixture of such solvents. Preferred water-miscible organic solvents comprise any of those described above and mixtures thereof.

**[0045]** Preferred water-immiscible solvents comprise aliphatic hydrocarbons; esters (for example ethyl acetate) chlorinated hydrocarbons (for example dichloromethane), ethers (for example diethyl ether) and mixtures thereof.

**[0046]** When the liquid medium comprises a water-immiscible organic solvent, preferably it comprises a polar solvent (for example a  $C_{1-4}$ alkanol) to enhance the solubility of the dye in the liquid medium. It is especially preferred that where the liquid medium is an organic solvent free from water it comprises a ketone (especially methyl ethyl ketone) and/or an alcohol (especially a  $C_{1-4}$ alkanol, more especially ethanol or propanol).

**[0047]** The organic solvent free from water may be a single organic solvent or a mixture of two or more organic solvents. It is preferred that when the medium is an organic solvent free from water it is a mixture of 2 to 5 different organic solvents. This allows a medium to be selected which gives good control over the drying characteristics and storage stability of the ink.

**[0048]** Ink media comprising organic solvent free from water are particularly useful where fast drying times are required and particularly when printing onto hydrophobic and non-absorbent substrates, for example plastics, metal and glass.

**[0049]** Preferred low melting solid media have a melting point in the range from 60°C to 125°C. Suitable low melting point solids include long chain fatty acids or alcohols, preferably those with  $C_{18-24}$ chains, and sulphonamides. The compound of Formula (1) may be dissolved in the low melting point solid or may be finely dispersed in it.

[0050] The ink may also contain additional components conventionally used in inks for IJP, for example viscosity and surface tension modifiers, corrosion inhibitors, biocides, migration reducing additives and surfactants which may be ionic or non-ionic.

5 [0051] A further aspect of the invention provides a process for printing an image on a substrate comprising applying to the substrate by means of an ink jet printer, an ink of the present invention as defined herein.

[0052] The ink jet printer preferably applies the ink to the substrate in the form of droplets which are ejected through a small orifice onto the substrate. Preferred ink jet printers are piezoelectric ink jet printers and thermal ink jet printers. In thermal ink jet printers, programmed pulses of heat are applied to the ink in a reservoir (e.g. by means of a resistor adjacent to the orifice) thereby causing the ink to be ejected in the form of small droplets directed towards the substrate 10 during relative movement between the substrate and the orifice. In piezoelectric ink jet printers the oscillation of a small crystal causes ejection of the ink from the orifice.

[0053] A further aspect of the present invention provides a substrate which has applied thereon an ink of the present invention as defined herein and/or one or more compounds of the present invention as defined herein.

15 [0054] The substrate preferably comprises paper, plastic, a textile, metal or glass, more preferably paper, an overhead projector slide or a textile material, especially paper. Preferred papers are plain or treated papers which may have an acid, alkaline or neutral character.

[0055] Preferably the ink and/or compounds of the present invention have been applied to the substrate by a printed process, more preferably the process of the present invention as defined herein.

[0056] When the substrate is a textile material the ink according to the invention is preferably applied thereto by:

20 i) applying the ink to the textile material using an ink jet printer; and  
ii) heating the printed textile material at a suitable temperature, preferably from 50°C to 250°C, to fix the ink on the material.

25 [0057] Preferred textile materials are natural, synthetic and semi-synthetic materials. Examples of preferred natural textile materials include wool, silk, hair and cellulosic materials, particularly cotton, jute, hemp, flax and linen. Examples of preferred synthetic and semi-synthetic materials include polyamides, polyesters, polyacrylonitriles and polyurethanes.

30 [0058] Preferably the textile material has been treated with an aqueous pre-treatment composition comprising a thickening agent and optionally a water-soluble base and a hydrotropic agent and dried prior to step i) above.

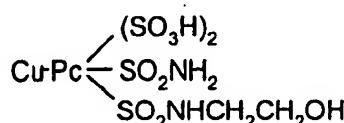
[0059] The pre-treatment composition preferably comprises a solution of the base and the hydrotropic agent in water containing the thickening agent. Particularly preferred pre-treatment compositions are described more fully in EP 0534660-A.

35 [0060] The invention is further illustrated by the following Examples in which all parts and percentages are by weight unless otherwise stated.

#### Example 1

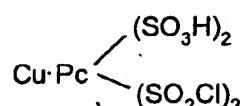
[0061] Preparation of

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a) Preparation of

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Copper phthalocyanine (115g) was added in portions to stirred chlorosulphonic acid (308 ml) over 30 minutes keeping the temperature below 50°C. The mixture was stirred for 30 minutes before being heated gradually to 140°C, at which temperature it was stirred for 3 hours to obtain CuPc(SO<sub>3</sub>H)<sub>4</sub>.

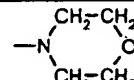
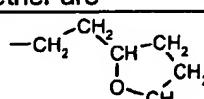
The mixture was cooled to 40°C and phosphorus trichloride (26.3g) was added in portions over 30 minutes, whilst the temperature was kept below 50°C. The mixture was stirred overnight at room temperature and then cooled to 0°C and poured onto a mixture of ice (700g), water (700g), concentrated hydrochloric acid (40 ml) and sodium chloride (100g). The mixture was stirred at 0°C for 30 minutes. The precipitated product was collected by filtration at reduced pressure and washed with ice-cold hydrochloric acid solution (0.5 M, 1.5 l) to obtain a paste of the above sulphonyl chloride.

(b) Preparation of title dye

Water (2.8 l), ethanolamine (16.6 g) and ammonia solution (48.6 g) were mixed in a 5 litre beaker and then placed in an ice bath (pH = 11.53, temperature 8° C). Sulphonyl chloride paste (862 g, prepared as described in step 1a above) was added to the mixture over 5 minutes and the reaction temperature rose to 10° C. Sodium hydroxide (165 mls 10% v/v) was added over 40 minutes to adjust the pH to 8.03. More NaOH (500 mls) was added to the mixture in aliquots over 30 minutes followed by a further 35 mls of NaOH to adjust the pH to 8.62. The pH of the mixture was maintained at 8.5 and the reaction mixture was stirred for five days. Sodium chloride (25% w/v) was added and the pH of the mixture was adjusted to 0.5 with concentrated hydrochloric acid. The resultant precipitate was collected by filtration at reduced pressure to obtain as the product the title dye which was characterised by HPLC and LC-MS (m/z 937, 873).

### Examples 2 to 15

**[0062]** The following examples were prepared analogously to Example 1, above replacing the ethanolamine in step 1(c) above with the molar equivalent of the appropriate amine of formula  $\text{NHR}^1\text{R}^2$ .

Example	$-R^1$	$-R^2$
2	$-C_2H_4OH$	$-C_2H_4OH$
3	$-H$	$-C_2H_4NHCH_2CO_2H$
4	$-H$	$-C_2H_4NHC_2H_4SO_3H$
5	$-H$	$-C_2H_4NHC_{16}H_{33}$
6	$R^1, R^2 \text{ & N together are}$ 	
7	$-H$	
8	$-H$	$-C_2H_4NHC_2H_4OH$
9	$-H$	$-C_2H_4NHC_{12}H_{25}$
10	$-H$	$-C_2H_4NHC_7H_{15}$
11	$-H$	$-C_2H_4NHC(CH_3)_2CH_2C(CH_3)_3$
12	$-H$	$-C_2H_4NHC_{10}H_{21}$
13	$-H$	$-C_2H_4NHC_{14}H_{29}$
14	$-H$	$-C_2H_4NHC_2H_4C(CH_3)_2C(CH_3)_3$
15	$-H$	$-C_2H_4SC_{10}H_{21}$

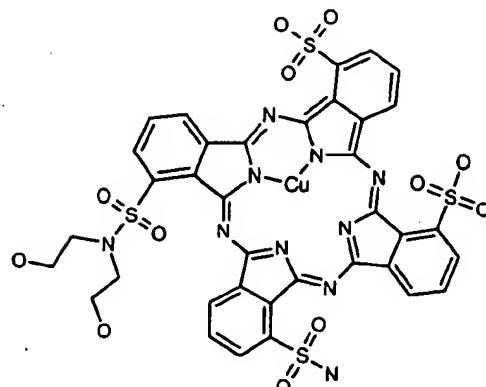
Alternative preparationsExample 16 (alternative preparation of Example 2)

5 [0063]

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Copper phthalocyanine (11.52g @ 98%) was added to chlorosulphonic acid (31.2ml) whilst maintaining the temperature < 60°C. The mixture was then heated to 140°C and kept at this temperature for ~3hours. The mixture was then cooled to 40°C and  $\text{PCL}_3$  (5.22g) was added dropwise, keeping the temperature at ~40°C. The mixture was then heated to 90°C for 2 hours, and then cooled to 0°C and added to a mixture of water (70g), ice (70g), salt (10g) and concentrated HCl (4ml). The precipitate obtained was then collected by filtration and washed with dilute acid and dried under reduced pressure. The resultant solid was added to a solution of diethanolamine (3.15g) in water (100ml) and the pH was adjusted to 8.5 by addition of 2N ammonium hydroxide solution. The mixture was stirred for one hour at 40°C at a pH from 8.0 to 8.5 and then at room temperature overnight. The mixture was then heated to 40°C and salted to 12% w/v with sodium chloride. The pH of the mixture was reduced to 0.6 by adding concentrated HCl. The solid obtained was collected by filtration and washed with brine. The solid was re-dissolved in distilled water at pH 7.5 and desalinated by dialysis. Water was evaporated to yield 17.4g of the title compound.

Example 17 (alternative preparation of Example 6)

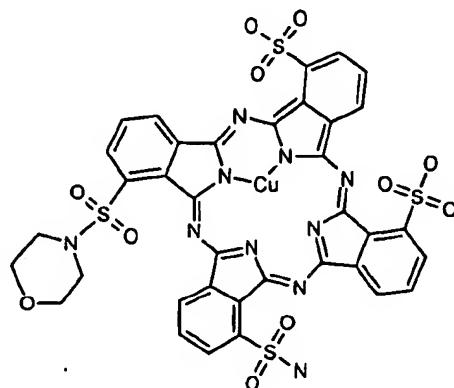
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[0064]

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17(a) Copper phthalocyanine (117.6g @ 98%) was added to chlorosulphonic acid (337 ml) over ~ 1 hour maintaining the temperature < 60°C throughout. The mixture was stirred for 15 min. then heated to 140°C over ~ 1.5 hours and stirred at this temperature for a further 3 hours. The mixture was cooled to 40°C and  $\text{PCL}_3$  (28.1g) was added dropwise over 1.3 hours at 40 to 50°C. The mixture was heated to 90°C for 3 hours and then cooled to room temperature and added to a mixture of salt (80g), water (600ml), ice (3kg) and concentrated HCl (30mls). The temperature of the mixture was kept below 5°C by addition of ice where necessary. The solid precipitate

obtained was collected by filtration and washed well with ice cold brine (1 litre) and dried under reduced pressure to obtain a filter cake (760.6g) which was used in the next step and Example 18.

5 17(b) Morpholine (14.2 g) was added to distilled water (320ml) followed by sufficient internal ice to lower temp to ~0°C. Filter cake [380.3 g - prepared as described in Example 17(a) above] was added to the mixture and the pH adjusted to 8.5 to 9.0 using ammonia solution (SG 0.88). The mixture was stirred at this pH and ~0°C for 4 hours and then heated to 40°C and held at pH 7.5 to 8.0 for 20 minutes. Concentrated HCl was then added to the mixture to lower its pH to ~1. The mixture was stirred for 30 minutes at this pH. The precipitate obtained was collected by filtration and washed with 2% HCl. The resultant paste was re-dissolved in distilled water (~2 l) at pH 9 to obtain a solution which was desalinated by reverse osmosis. Water was evaporated from this solution to yield 63g of the title compound. The respective ratio of  $\text{SO}_3\text{H}$  /  $\text{SO}_2\text{NH}_2$  /  $\text{SO}_2\text{N}(\text{C}_2\text{H}_2)_2\text{O}$  was determined to be 1.3 / 1.2 / 1.5.

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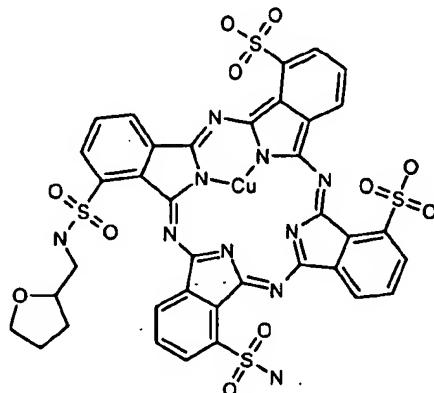
Example 18 (alternative preparation of Example 7)

15 [0065]

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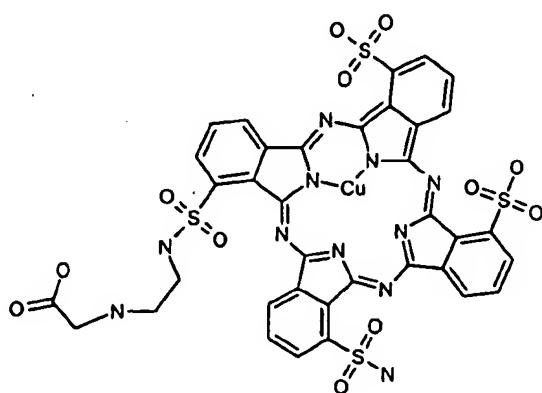
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Example 19 (alternative preparation of Example 3)

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[0068] Copper phthalocyanine (117.6g) was added to chlorosulphonic acid (307ml, 542.9g) over 1-2 hours whilst maintaining the temperature at < 60°C. The temperature was raised to 140°C over 1.5 hours and heated at that temperature for a further 3 hours. The mixture was cooled to 40°C and  $\text{PCL}_3$  (18ml, 27.5g) was added dropwise maintaining the temperature at ~40°C. The mixture was then heated to ~90°C for ~3 hours and then cooled to 40°C. This mixture was then added to a mixture of water (500ml), ice (2kg), concentrated HCl (20 ml) and salt (50g) keeping the temperature at ~0°C by addition of internal ice as necessary. The solid precipitate obtained was then collected by filtration and washed with dilute acid. This solid was then added to a mixture of iced-water (1.1 kg) and 2-chloroethylamine (46.8g) and the pH raised to 9 by addition of 10% ammonia solution. The mixture was then stirred at pH 8 to 9 and 0°C for 4 hours, after which the temperature was raised to 40°C and the mixture was stirred for 30 minutes. Concentrated HCl was added to lower the pH to 1 to 2. The solid precipitate obtained was then collected by filtration, washed with dilute acid to give (389 g) of solid which was used directly in the next step below.

[0069] The solid prepared above (389g) was added to water (1 litre) followed by glycine (20.26 g) and the pH of the mixture was adjusted to ~9 by addition of caustic liquor. The mixture was stirred at this pH and heated to 70 to 80°C for ~4 hours. Then 20% w/v salt was added to the mixture and the pH adjusted to 4 by addition of concentrated HCl.

[0070] The precipitate obtained was then collected by filtration, washed well with brine and then re-dissolved in distilled water at pH 9 to obtain a solution which was de-salinated by reverse osmosis. Water removed to yield, as solid, 56g of the title compound. The respective ratio of  $\text{SO}_3\text{H}$  /  $\text{SO}_2\text{NH}_2$  /  $\text{SO}_2\text{NHC}_2\text{H}_4\text{NHCH}_2\text{CO}_2\text{H}$  was determined to be 1.3 / 1.2 / 1.5

#### Further, alternative preparations

[0070] Examples 4 to 5 and 8 to 15 herein can all also be prepared by an analogous route to that described in Example 19 above by replacing the glycine with an equivalent molar amount of the corresponding amine or thiol.

[0071] An alternative route to the preparation of Example 1 above would be to remove the glycine stated above and heat at pH 10 to 12 and 90°C for 4 hours which results in hydrolysis of the Cl substituent to OH.

#### Results

##### Chroma

[0072] Example 1 herein was compared with the known copper phthalocyanine dye Cyan 1 [=  $\text{CuPc}(\text{SO}_3\text{H})_3(\text{SO}_2\text{NH}_2)_3$ ] in a conventional ink formulation. The chroma of an IJ print made with an ink of Example 1 was found to be 5% higher at a similar optical density than a comparative print made with Cyan 1 (the optical density of a print, can be conveniently measured using an X-Rite 938 Spectrodensitometer). Chroma values were obtained in a conventional manner.

Dye	Chroma
Example 1	40.5
Cyan 1 (known dye)	38.5

[0073] Examples 2, 6 and 7 were also tested for chroma and performed very similarly to Example 1, (i.e. better than Cyan 1).

##### Water fastness

[0074] Dyes of the present invention also demonstrate a desirable combination of solubility and water fastness compared to prior art dye, as illustrated by the following test.

[0075] The water fastness of Example 6 herein was compared with the known dye Cyan 1 in the conventional run down test. Water fastness (denoted herein by WF) was determined by running water (0.5 ml) down lines of print at an angle of approximately 45°C immediately after the lines had been printed. The prints were given a score of 1 to 10 where 1 indicates poor wet fastness and 10 indicates no detected ink run down (i.e. 100% water fast).

Dye	WF
Example 6	6
Cyan 1 (known dye)	3

[0076] Examples 5, 6, 9 to 15 herein were also tested in the run down test and showed a comparable WF to Example 6 (i.e. better than Cyan 1).

[0077] The aqueous solubility of Example 1 was found to be 29%. This compares to an aqueous solubility of 14% for its mono-sulphonic acid substituted analogue.

5 [0078] Thus contrary to what one would expect, the applicant has found that improved water solubility in the dyes of the present invention (which equates to greater operability in IJP) does not occur at the expense of poor WF.

[0079] Prints made with the dyes of the Examples exhibit a WF comparable to or better than the corresponding zero or mono-sulphonic acid dyes. Dyes of the invention also exhibit a particularly attractive cyan shade with good chroma compared to their corresponding zero or mono-sulphonic acid analogues.

10 [0080] The dyes of the present invention are significantly easier than prior art dyes to formulate in IJP inks and use in IJP and yet they produce prints of comparable or improved quality to prior art dyes.

#### Salts

15 [0081] The exemplified sodium salts (examples 1 to 19) prepared as described above, may be used after suitable purification directly in ink formulations as described below or may be converted to other IJP-effective salts as follows. Each example was dissolved in distilled water. The solution was filtered and the volume of the filtrate adjusted to 4 litres. The solution was desalinated by reverse osmosis. The dissolved sodium salt may be converted into other salt [e.g. the potassium salt] by passing the solution through an ion-exchange column comprising Dowex HGRW resin 20 saturated with a suitable solution [e.g. potassium hydroxide solution (5% w/v)]. The solution of the (e.g. potassium) salt thus obtained was then filtered and water was evaporated from the filtrate to yield a purified salt suitable for use directly in an ink as described below.

#### Inks

25 [0082] The effectiveness in ink jet printing of compounds of Formula (1) was demonstrated as follows. Inks separately comprising each exemplified dye were prepared by dissolving 2 parts of the sodium salt, prepared as described above, in 98 parts of a mixture of water and 2-pyrrolidone (in a respective ratio of 90:10 by volume). The inks were printed onto plain paper (obtained from Felix Schoeller) using a thermal ink-jet printer to give a bright cyan prints which had 30 excellent light fastness.

[0083] Further inks comprising the exemplified dyes may be prepared as described in the following tables in which the number in the first column (headed Ex. no.) denotes the example number of dye to be used in the ink. The dye may be in its free acid form and/or in the form of any IJP-effective salt (e.g. sodium, potassium, ammonium, or QAC salt). Numbers quoted in the second column onwards refer to the number of parts of the relevant ingredient 35 and all parts are by weight. The inks may be applied to paper by thermal or piezo ink jet printing.

[0084] The following abbreviations are used in the tables:

PG = propylene glycol;	DEG = diethylene glycol;	NMP = N-methyl pyrrolidone;
DMK = dimethylketone;	NaST = Na stearate	IPA = isopropanol;
MEOH = methanol;	2P = 2-pyrrolidone;	MIBK = methylisobutyl ketone;
CET = cetyl ammonium bromide;	TBT = tertiary butanol;	TDG = thioglycol;
BDL = butane-2,3-diol;	PHO = Na <sub>2</sub> HPO <sub>4</sub> ;	P12 = propane-1,2-diol.
CHL = cyclohexanol; and	PDL = pentan-1,5-diol.	

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TABLE I

Ex. no.	Dye	Water	PG	DEG	NMP	DMK	MEOH	2P	MIBK
1	2.0	80	5		6	4		5	
2	10.0	85	3		3	3	5	1	
3	2.1	91		8					1
4	2.4	75	3	4		5	6		5
5	5.1	96					4		
6	1.8	80		5				15	

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TABLE I (continued)

Ex. no.	Dye	Water	PG	DEG	NMP	DMK	MEOH	2P	MIBK
7	2.6	84			11			5	
8	3.3	80	2			10	2		6
9	5.4	69	2	20	2	1		3	3
10	7.0	70			15		3		10

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TABLE II

Ex. No.	Dye	Water	PG	DEG	NMP	NaOH	Na ST	IPA	2P	MIBK
11	2.0	70		7		6		3		2
12	4.0	65	4		6		0.7		5	
13	1.0	50		4				5		1
14	3.1	86	5			2	0.2	4		5
15	1.1	81		2	9	0.5	0.5		9	

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TABLE III

Ex. no.	Dye	Water	PG	DEG	NMP	DMK	IPA	MEOH	2P	MIBK
1	3.1	86	5				4			5
2	1.1	81			9				9	
3	2.5	60	4	15	3	3	6	10	5	4
4	3.2	65		5	4	6	5	4	6	5
5	10.0	80	2	6	2	5	1			4

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TABLE IV

Ex. no.	Dye	Water	PG	DEG	NMP	DMK	NaOH	Na ST	IPA
6	3.0	90		5	5		0.2		
7	5	65	5	20					10
8	4.1	80		5	2	10		0.3	
9	10.8	90	5						5
10	12.0	90				7	0.3		3

TABLE V

Ex. no.	Dye	Water	PG	DEG	NMP	CET	PHO	2P	P12
11	3.0	80	15			0.2		5	
12	9.0	90		5			1.2		5
13	2.5	90		6	4		0.12		
14	3.1	82	4	8		0.3			6
15	10.0	91			6			3	
1	5.0	78	5	11				6	

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TABLE V (continued)

Ex. no.	Dye	Water	PG	DEG	NMP	CET	PHO	2P	P12
2	6.0	63	3		4		2.0		
3	3.0	72		15		0.8			3
4	5.4	86	5		7		3.0	7	
5	2.0	90		10					10

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TABLE VI

Ex. no.	Dye	Water	PG	DEG	NMP	CET	TBT	TDG
6	1.5	85	5	5		0.15	5.0	0.2
7	9.0	90		5	5			0.3
8	2.0	90		10				
9	2.0	88						10
10	5.5	70	4		4	0.4	3	

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TABLE VII

Ex. no.	Dye	Water	PG	DEG	NMP	TBT	TDG	BDL	PHO	2P
11	0.9	85		10				5	0.2	
12	4.0	70		10	4			1		4
13	2.2	75	4	10	3			2		6
14	9.0	76		9	7	3.0			0.95	5
15	2.1	70	5	5	5	0.2	0.1	5	0.1	5

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TABLE VIII

Ex. no.	Dye	Water	PG	DEG	NMP	TBT	TDG	2P
1	3.0	55			5	2.0		3
2	6.0	65		4		0.1	5	
3	5.0	78			5		12	5
4	8.0	70	2		8		15	5
5	10.0	80					8	12

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TABLE IX

Ex. no.	Dye	Water	PG	DEG	NMP	BDL	PHO	2P	P12
6	10.0	75	3		5	3		3	1
7	3.5	80		6					5
8	2.0	90	7			7	0.5		
9	6.0	65			5			2	
10	4.0	70		10	4	1		4	11

TABLE X

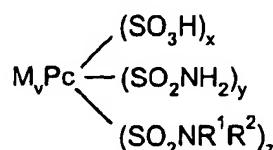
Ex. no.	Dye	Water	PG	DEG	NMP	CET	TBT	TDG	BDL	PHO	2P	
5	11	1.5	80	1	2	3	0.5	0.4		7	2	10
10	12	3.0	60		4		2.0		0.5			
15	13	4.5	90		3	6		7.0		1	3	2
20	14	3.0	95		7		4		3		0.5	
	15	2.1	70	5	5	5	0.1	0.2	0.1	5	0.1	5

TABLE XI

Ex. no.	Dye	Water	PG	DEG	NMP	CHL	PHO	2P	PDL
15	1	7.0	75	3		5	3		3 1
20	2	3.5	60			2			5
25	3	2.0	90	7			7	1.5	2
	4	6.0	65			5			2
	5	4.0	70		5	4	1		4 12

## 25 Claims

1. One or more compounds of Formula (1) and salts thereof:



35 Formula (1)

in which:

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M represents a metal or H;

Pc represents a phthalocyanine nucleus; and

45 R<sup>1</sup> represents H or -(CH<sub>2</sub>)<sub>n</sub>R<sup>3</sup>; R<sup>2</sup> represents -(CH<sub>2</sub>)<sub>n</sub>R<sup>3</sup>; or R<sup>1</sup> and R<sup>2</sup> together with the nitrogen atom to which they are attached represent a 5 or 6-membered ring; where:

n is independently an integer from 1 to 30; and

50 R<sup>3</sup> is independently a group selected from H, -SR<sup>4</sup>, -CO<sub>2</sub>R<sup>5</sup> and -NR<sup>6</sup>R<sup>7</sup>; where:

R<sup>4</sup>, R<sup>5</sup>, R<sup>6</sup> and R<sup>7</sup> independently represent H, C<sub>1-30</sub>alkyl optionally substituted by one or more groups selected from hydroxy, mercapto, sulpho, carboxy, cyano and -PO<sub>3</sub>H<sub>2</sub>;

v is the valence of Pc divided by the valence of M;

x is from 1.8 to 3.8;

y is from 0.1 to 2.7;

z is from 0.1 to 2.7; and

x, y and z satisfy 2 ≤ x + y + z ≤ 4;

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with the provisos that:

R<sup>4</sup> and R<sup>5</sup> are other than both H;

when R<sup>1</sup> is H or C<sub>1-4</sub>alkyl then R<sup>4</sup> is other than C<sub>1-4</sub>alkyleneSO<sub>3</sub>H;

when R<sup>1</sup> is H or C<sub>1-4</sub>alkyl and R<sup>6</sup> is H, C<sub>1-4</sub>alkyl or hydroxyC<sub>1-4</sub>alkyl; then R<sup>7</sup> is other than C<sub>1-4</sub>alkyl, hydroxyC<sub>1-4</sub>alkyl or morpholino; and  
when R<sup>1</sup> is H or C<sub>1-4</sub>alkyl then R<sup>6</sup> and R<sup>7</sup> are other than both H or (together with the N atom to which they are attached) other than a morpholino ring.

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2. One or more compounds according to claim 1 wherein M is Cu or Ni, R<sup>1</sup> is selected from H and optionally substituted C<sub>1-15</sub>alkyl and R<sup>2</sup> is a substituted C<sub>1-15</sub>alkyl.
3. One or more compounds according to any one of the preceding claims wherein M is Cu, R<sup>1</sup> is selected from H, C<sub>1-4</sub>alkyl and hydroxyC<sub>1-4</sub>alkyl and R<sup>2</sup> is a hydroxyC<sub>1-4</sub>alkyl.
- 10 4. One or more compounds according to any one of the preceding claims wherein x is 1.8 to 2.2.
- 15 5. One or more compounds according to any one of the preceding claims wherein y and z are independently from 0.5 to 1.5.
6. One or more compounds according to any one of the preceding claims wherein x is 2 and y and z are each 1.
- 20 7. An ink which is effective for use in ink jet printing, the ink comprising a fluid medium and a colorant comprising one or more compounds of Formula (1) according to any one of the preceding claims.
8. A process for printing an image on a substrate comprising applying to the substrate by means of an ink jet printer, an ink as claimed in claim 7.
- 25 9. A substrate which has applied thereon, an ink as claimed in claim 7 and/or one or more compounds according to any one of claims 1 to 6.

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DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
Y	EP 0 719 847 A (SEIKO EPSON CORP) 3 July 1996 (1996-07-03) * abstract; examples *	1-9	C09B47/26 C09D11/00
Y	EP 0 196 901 A (TAOKA CHEMICAL CO LTD ;SUMITOMO CHEMICAL CO (JP)) 8 October 1986 (1986-10-08) * abstract; example 4 *	1-9	
Y	EP 0 596 383 A (BAYER AG) 11 May 1994 (1994-05-11) * abstract; examples *	1-9	
Y	WO 97 13811 A (KENYON RONALD WYNFORD ;ZENECA LTD (GB); MISTRY PRAHALAD MANIBHAI ) 17 April 1997 (1997-04-17) * abstract *	1-6	
Y	EP 0 519 395 A (EASTMAN KODAK CO) 23 December 1992 (1992-12-23) * abstract; example 33 *	1-9	
Y	EP 0 649 881 A (BAYER AG) 26 April 1995 (1995-04-26) * abstract; examples 7,8 *	1-9	C09B C09D
P,Y	WO 98 49239 A (CARR KATHRYN ;ZENECA LTD (GB)) 5 November 1998 (1998-11-05) * abstract; examples 2-5 *	1-9	
A	FR 1 347 692 A (DURAND & HUGUENIN) 4 January 1964 (1964-01-04) * example 3 *	1-9	
P,A	WO 98 49240 A (CARR KATHRYN ;ZENECA LTD (GB)) 5 November 1998 (1998-11-05) * abstract *	1-9	
		-/-	
The present search report has been drawn up for all claims			
Place of search		Date of completion of the search	Examiner
THE HAGUE		9 July 2002	Dauksch, H
CATEGORY OF CITED DOCUMENTS			
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document			
T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons B : member of the same patent family, corresponding document			



DOCUMENTS CONSIDERED TO BE RELEVANT		Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)		
Category	Citation of document with indication, where appropriate, of relevant passages				
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